In the Claims:

- 1. (Original) A bivalent or multivalent antibody characterized by the following features:
 - (a) it is capable of supressing an immune reaction;
 - (b) it is devoid of constant antibody regions; and
 - (c) it binds an epitope on the CD3 complex of the T-cell receptor.
- 2. (Original) The antibody of claim 1 that is a diabody.
- 3. (Original) The antibody of claim 1 that comprises two scFv antibodies linked by a peptide linker.
- 4. (Original) The antibody of claim 1 that is a single chain diabody.
- 5. (Currently Amended) The antibody according to any one of claims 1 to 4 claim 1, wherein the its variable V_H and V_L domains are connected via the peptide linker SAKTTP or SAKTTPKLGG.
- 6. (Currently Amended) The antibody according to any one of claims 1 to 5 claim 1, wherein the its variable domains correspond to the variable domains of the an antibody produced by the hybridoma of ATCC deposit number CRL 8001.
- 7. (Currently Amended) The antibody according to claim 6, wherein a cysteine at position H100A (Kabat numbering system) has been exchanged for another amino acid.
- 8. (Original) The antibody according to claim 7, wherein the cysteine has been exchanged for a serine.

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- 9. (Currently Amended) A polynucleotide, which encodes an antibody of any one of claims 1 to 8 according to claim 1.
- 10. (Original) An expression vector comprising the polynucleotide of claim 9.
- 11. (Original) The expression vector of claim 10, which is pSKK3-scFv_6-anti-CD3 (DSM 15137).
- 12. (Original) A host cell containing the expression vector of claim 10 or 11.
- 13. (Currently Amended) A pharmaceutical composition eentaining comprising the antibody of any one of claims 1 to 8 claim 1, the polynucleotide of claim 9, or the expression vector of claim 10 or 11.
- 14. (Currently Amended) Use of an antibody which is characterized by the following features:
 - (a) it is capable of supressing an immune reaction;
 - (b) it is devoid of constant antibody regions; and
 - (c) it binds an epitope on the CD3 complex of the T cell receptor; or the polynucleotide of claim 9 or the expression vector of claim 10 or 11 for the preparation of a pharmaceutical composition A method for immunotherapy comprising the step of administering to a subject the pharmaceutical composition according to claim 13.
- 15. (Currently Amended) Use according to claim 14, wherein the antibody is A method for immunotherapy comprising the step of administering to a subject a pharmaceutical composition comprising the antibody of any one of claims 1 to 8 claim 1.

- 16. (Currently Amended) Use The method according to claim 14 or 15, wherein said immunotherapy is a therapy against acute transplant rejections.
- 17. (Currently Amended) Use—of A method for gene therapy comprising the step of administering to a subject a pharmaceutical composition comprising the polynucleotide of claim 9 or the expression vector of claim 10 or 11 for the preparation of a pharmaceutical composition for gene therapy.